What is claimed is:

1. A method of changing a gynecological condition of a female comprising:

evaluating the condition of a uterus of said female;

introducing a presterilized implant into said uterus;

contacting said implant with uterine tissue so as to induce a tissue response in said uterus;

maintaining contact between said implant and said uterine tissue for at least so long that said tissue response causes a changed gynecological condition in said female.

- 2. A method according to claim 1, wherein the contact between said implant and said uterine tissue is maintained at least until adhesions are formed in said uterus.
- 3. A method according to claim 1, wherein the contact between said implant and said uterine tissue is maintained at least until contraception in said uterus is achieved.
- 4. A method according to claim 1, wherein the contact between said implant and said uterine tissue is maintained at least until menorragia has been substantially eliminated in said female.
- 5. A method according to claim 1, wherein the contact between said implant and said uterine tissue is maintained at least until walls of said uterus adhere together.
- 6. A method according to claim 1, wherein said presterilized implant is coated with an adhesion inducing substance.
- 7. A method according to claim 6, wherein said presterilized implant is coated with a biologic coating prior to introducing said implant into said uterus.
- 8. A method according to claim 1, wherein said presterilized implant is formulated at least in part from polyester prior to its introduction into said uterus.

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 9. A method according
 - 9. A method according to claim 1, wherein said presterilized implant is introduced through a catheter.
 - 10. An implant for changing the gynecological state of a female comprising: a presterilized substance;

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said substance configured for causing a tissue response in uterine tissue; and,

said substance sized and shaped for sufficiently contacting uterine tissue such that said tissue response causes a gynecological change in said female.

- 11. An implant according to claim 10, wherein said presterilized substance is a mesh material.
- 12. An implant according to claim 10, wherein said presterilized substance is a polyester material.
- 13. An implant according to claim 10, wherein said presterilized substance is coated with an adhesion inducing substance.
- 14. An implant according to claim 10, wherein said presterilized substance includes a frame, at least a portion of which is covered by a mesh material.
- 15. An implant according to claim 14, wherein said mesh material is comprised substantially of polyester.
- 16. An implant according to claim 15, wherein said frame includes a plurality of linear extensions, at least two of such extensions sized and shaped to extend across a uterine wall.
- 17. An implant according to claim 16, wherein said at least two extensions are movable between a collapsible and a deployed position.
- 18. An implant according to claim 10, wherein said substance is sized and shaped so as to eliminate menorragia.
- 19. An implant according to claim 10, wherein said substance is sized and shaped so as to cause contraception in said uterus.